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## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims

## 1. (previously presented) A compound of the Formula A:

$$(R^1)_{\widehat{n}} \xrightarrow{N} (R^2)_{\widehat{n}} R^4$$

wherein:

a is 0 or 1; b is 0 or 1; m is 0, 1 or 2; n is 0, 1, 2 or 3; p is 0, 1 or 2; q is 0, 1, 2 or 3; r is 0 or 1; s is 0 or 1; t is 2, 3, 4, 5 or 6;



is heterocyclyl;

Q is pyrimidinyl pyrazole optionally substituted with one to three Rz;

 $R^{1} \ is \ independently \ selected \ from: 1) \ (C=O)_{a}O_{b}C_{1}-C_{10} \ alkyl, 2) \ (C=O)_{a}O_{b}aryl, 3) \ C_{2}-C_{10} \ alkenyl, 4) \ C_{2}-C_{10} \ alkynyl, 5) \ (C=O)_{a}O_{b} \ heterocyclyl, 6) \ (C=O)_{a}O_{b}C_{3}-C_{8} \ cycloalkyl, 7) \ (CO_{2}H, 8) \ halo, 9) \ CN, 10) \ OH, 11) \ O_{b}C_{1}-C_{6} \ perfluoroalkyl, 12) \ O_{a}(C=O)_{b}NR67, 13) \ NRe(C=O)_{b}NR687, 14) \ S(O)_{m}R^{a}, 15) \ S(O)_{2}NR687, 16) \ NReS(O)_{m}R^{a}, 17) \ oxo, 18) \ CHO, 19) \ NO_{2}, 20) \ NRe(C=O)_{O}b_{R}^{a}, 21) \ O(C=O)O_{b}C_{1}-C_{10} \ alkyl, 22) \ O(C=O)O_{b}C_{3}-C_{8} \ cycloalkyl, 23) \ O(C=O)O_{b}aryl, 24) \ C_{1}-C_{6}alkyl(C=NRb)N(R^{b})_{2}, 25) \ O(C=O)O_{b}-heterocycle, 26) \ O_{a}^{a}-P=O(OH)_{2} \ and 27) \ -N=CHN(R^{b})_{2}, \ said \ alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one or more substituents selected from <math display="inline">R^{z}$ ;

 $R^2$  is independently selected from: 1) (C=O)<sub>a</sub>O<sub>b</sub>C<sub>1</sub>-C<sub>10</sub> alkyl, 2) (C=O)<sub>a</sub>O<sub>b</sub>aryl, 3) C<sub>2</sub>-C<sub>10</sub> alkenyl, 4) C<sub>2</sub>-C<sub>10</sub> alkynyl, 5) (C=O)<sub>a</sub>O<sub>b</sub> heterocyclyl, 6) (C=O)<sub>a</sub>O<sub>b</sub>C<sub>3</sub>-C<sub>8</sub> cycloalkyl, 7) CO<sub>2</sub>H, 8) halo, 9) CN,

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10) OH, 11)  $O_bC_1$ - $C_6$  perfluoroalkyl, 12)  $O_a(C=O)_bNR6R^7$ , 13)  $NRe(C=O)NR6R^7$ , 14)  $S(O)_mR^a$ , 15)  $S(O)_2NR6R^7$ , 16)  $NReS(O)_mR^a$ , 17) CHO, 18)  $NO_2$ , 19)  $NRe(C=O)O_bR^a$ , 20)  $O(C=O)O_bC_1$ - $C_{10}$  alkyl, 21)  $O(C=O)O_bC_3$ - $C_8$  cycloalkyl, 22)  $O(C=O)O_b$ -aryl, 23)  $O(C=O)O_b$ -heterocycle, and 24)  $O_a$ - $O(OH)_2$ , said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one, two or three substituents selected from  $R^z$ ;

R3 and R4 are independently selected from: H, C1-C6-alkyl and C1-C6-perfluoroalkyl, or

R<sup>3</sup> and R<sup>4</sup> are combined to form -(CH<sub>2</sub>)<sub>t</sub>- wherein one of the carbon atoms is optionally replaced by a moiety selected from O, S(O)<sub>m</sub>, -N(Rb)C(O)-, and -N(COR<sup>a</sup>)-;

 $R^5$  is independently selected from: 1) (C=O)\_aOb\_C1-C1\_0 alkyl, 2) (C=O)\_aOb\_aryl, 3) C2-C1\_0 alkenyl, 4) C2-C1\_0 alkynyl, 5) (C=O)\_aOb heterocyclyl, 6) (C=O)\_aOb\_C3-C8 cycloalkyl, 7) CO2H, 8) halo, 9) CN, 10) OH, 11) Ob\_C1-C6 perfluoroalkyl, 12) Oa(C=O)bNR6R7, 13) NRc(C=O)NR6R7, 14) S(O)\_mRa, 15) S(O)\_2NR6R7, 16) NRcS(O)\_mRa, 17) oxo, 18) CHO, 19) NO2, 20) O(C=O)Ob\_C1-C1\_0 alkyl, 21) O(C=O)Ob\_C3-C8 cycloalkyl, and 22) Oa-P=O(OH)2, said alkyl, aryl, alkenyl, alkynyl, heterocyclyl, and cycloalkyl optionally substituted with one or more substituents selected from  $\mathbb{R}^z;$ 

R6 and R7 are independently selected from: 1) H, 2) (C=O)O<sub>b</sub>Ra, 3) C1-C1<sub>0</sub> alkyl, 4) aryl, 5) C2-C1<sub>0</sub> alkenyl, 6) C2-C1<sub>0</sub> alkynyl, 7) heterocyclyl, 8) C3-C8 cycloalkyl, 9) SO<sub>2</sub>Ra, 10) (C=O)NRb<sub>2</sub>, 11) OH, and 12) O<sub>a</sub>-P=O(OH)<sub>2</sub>, said alkyl, cycloalkyl, aryl, heterocylyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from RZ or

R6 and R7 can be taken together with the nitrogen to which they are attached to form a monocyclic or bicyclic heterocycle with 4-7 members in each ring and optionally containing, in addition to the nitrogen, one or more additional heteroatoms selected from N, O and S, said monocyclic or bicyclic heterocycle optionally substituted with one or more substituents selected from Rz;

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heterocyclyl is optionally substituted with up to three substituents selected from Rb, OH, (C1-C6)alkoxy, halogen, CO2H, CN, O(C=O)C1-C6 alkyl, oxo, N(Rb)2 and Oa-P=O(OH)2;

R<sup>a</sup> is: substituted or unsubstituted (C<sub>1</sub>-C<sub>6</sub>)alkyl, substituted or unsubstituted (C<sub>2</sub>-C<sub>6</sub>)alkenyl, substituted or unsubstituted (C<sub>2</sub>-C<sub>6</sub>)alkynyl, substituted or unsubstituted (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, substituted or unsubstituted aryl, (C<sub>1</sub>-C<sub>6</sub>)perfluoroalkyl, 2,2,2-trifluoroethyl, or substituted or unsubstituted heterocyclyl; and

Rb is: H, (C<sub>1</sub>-C<sub>6</sub>)alkyl, substituted or unsubstituted aryl, substituted or unsubstituted benzyl, substituted or unsubstituted heterocyclyl, (C<sub>3</sub>-C<sub>6</sub>)cycloalkyl, (C=O)OC<sub>1</sub>-C<sub>6</sub> alkyl, (C=O)C<sub>1</sub>-C<sub>6</sub> alkyl or S(O)<sub>2</sub>Ra;

Re is selected from: 1) H, 2) C1-C10 alkyl, 3) aryl, 4) C2-C10 alkenyl, 5) C2-C10 alkynyl, 6) heterocyclyl, 7) C3-C8 cycloalkyl, and 8) C1-C6 perfluoroalkyl, said alkyl, cycloalkyl, aryl, heterocylyl, alkenyl, and alkynyl is optionally substituted with one or more substituents selected from Rz, or or a pharmaceutically acceptable salt or a stereoisomer thereof.

(previously presented) The compound according to Claim 1 of the Formula B:

$$(R^1)_n \xrightarrow{N} \bigoplus_{B} (R^2)_{b}$$

or a pharmaceutically acceptable salt or a stereoisomer thereof.

(previously presented) The compound according to Claim 2 wherein:

Q is pyrimidinyl pyrazole optionally substituted with one to three RZ;

Ra is: (C1-C6)alkyl, (C3-C6)cycloalkyl, aryl, or heterocyclyl; and

 $R^b$  is: H,  $(C_1-C_6)$ alkyl, aryl, heterocyclyl,  $(C_3-C_6)$ cycloalkyl, (C=O)OC1-C6alkyl, (C=O)C1-C6alkyl or S(O)2 $R^a$ ;

or a pharmaceutically acceptable salt or a stereoisomer thereof.

4. (original) The compound according to Claim 3 wherein:

q is 0;

R<sup>2</sup> is independently selected from: 1) C<sub>1</sub>-C<sub>6</sub> alkyl, 2) aryl, 3) heterocyclyl, 4) CO<sub>2</sub>H, 5) halo, 6) CN, 7) OH, 8) S(O)<sub>2</sub>NR<sup>6</sup>R<sup>7</sup>, and 9) O<sub>a</sub>-P=O(OH)<sub>2</sub>, said alkyl, aryl and heterocyclyl optionally substituted with one, two or three substituents selected from Rz;

or a pharmaceutically acceptable salt or a stereoisomer thereof.

5. (previously presented) The compound according to Claim 4 of the Formula C:

$$(R^1)_n$$

wherein:

n is 0, 1 or 2:

Q is pyrimidinyl pyrazole optionally substituted with one to three Rz;

or a pharmaceutically acceptable salt or a stereoisomer thereof.

6. (previously presented) A compound which is selected from:

1-(1-{4-[5-(5-amino-1,3,4-thiadiazol-2-yl)-3-phenylpyridin-2-yl]benzyl}, piperidin-4-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine;

1-(1-{4-[5-(1,2,4-oxadiazol-3-yl)-3-phenylpyridin-2-yl]benzyl}piperidin-4-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine:

1-(1-{4-[3-phenyl-5-(1H-1,2,4-triazol-5-yl)pyridin-2-yl]benzyl}piperidin-4-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine;

1-{1-{4-(3-phenyl-5-pyrimidin-2-yl)pyridin-2-yl)benzyl]piperidin-4-yl}-1H-pyrazolo[3,4-d]pyrimidin-4-amine;

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1-{1-[4-(5'-phenyl-2,3'-bipyridin-6'-yl)benzyl]piperidin-4-yl}-1H-pyrazolo[3,4-d]pyrimidin-4-amine;

or a pharmaceutically acceptable salt or a stereoisomer thereof.

- 7-10. (canceled)
- 11. (original) A compound according to Claim 6 which is:

1-(1-{4-[5-(5-amino-1,3,4-thiadiazol-2-yl)-3-phenylpyridin-2-yl]benzyl}piperidin-4-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine:

or a pharmaceutically acceptable salt or a stereoisomer thereof.

- 12. (canceled)
- 13. (original) A compound according to Claim 6 which is:

1-(1-{4-[5-(1,2,4-oxadiazol-3-yl)-3-phenylpyridin-2-yl]benzyl}piperidin-4-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine:

or a pharmaceutically acceptable salt or a stereoisomer thereof.

14. (original) A compound according to Claim 6 which is:

 $1-\{1-\{4-(3-phenyl-5-pyrimidin-2-yl)ptridin-2-yl)ptridin-4-yl\}-1\\H-pyrazolo[3,4-d]pyrimidin-4-mine;$ 

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or a pharmaceutically acceptable salt or a stereoisomer thereof.

- 15. (original) A pharmaceutical composition comprising a pharmaceutical carrier, and dispersed therein, a therapeutically effective amount of a compound of Claim 1.
- 16. (original) A pharmaceutical composition comprising a pharmaceutical carrier, and dispersed therein, a therapeutically effective amount of a compound of Claim 6.
- 17. (currently amended) A method for treating eaneer <u>carcinoma</u> which comprises administering to a mammal in need thereof a therapeutically effective amount of a compound of Claim 1.
- 18. (currently amended) A method for treating eaneer <u>carcinoma</u> which comprises administering to a mammal in need thereof a therapeutically effective amount of a compound of Claim 6.

19-20. (canceled)